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File 351:Derwent WPI 1963-2001/UD, UM &UP=200211
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       Set Items Description
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WPI Acc No: 1989-160059/198922
Related WPI Acc No: 1988-028629
XRAM Acc No: C89-070912
   Quat derivs. of noroxymorphone - administered enterally before
   anaesthetic analgesic, for preventing nausea etc.
Patent Assignee: UNIV CHICAGO (UYCH-N)
. Inventor: GOLDBERG L I
Number of Countries: 004 Number of Patents: 006
Patent Family:
Patent No
              Kind
                     Date
                              Applicat No
                                             Kind Date
JP 1068376
               A 19890314
                              JP 87330356
                                                  19871228
                                                             198922 B
DK 8706933
                    19890304
                                                             198923
US 4861781
                   19890829
                              US 8792470
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CA 1315689
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                    19930406
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DK 167340
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JP 2625457
               B2 19970702 JP 87330356
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Priority Applications (No Type Date): US 8792470 A 19870903; US 86837399 A
   19860307
Patent Details:
Patent No Kind Lan Pg
                         Main IPC
                                      Filing Notes
JP 1068376
              A
                      4 P
DK 167340
              В
                                      Previous Publ. patent DK 8706933
JP 2625457
               B2
                                      Previous Publ. patent JP 1068376
DK 8706933
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US 4861781
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 CA 1315689
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Abstract (Basic): JP 1068376 A

Prevention and reduction of nausea and emesis caused from the usage of an anaesthetic analgesic to the homotherm; which comprises an administration of one or more of the compound (I) with an effective amount, before the administration of the anaesthetic analgesic or at the simultaneous use of the analgesic.

In (I) R = allyl or relative allyl such as chloroallyl, cyclopropyl-methyl or propargyl; X = acidic anion, esp. anion chloride, anion bromide, anion iodide or methylsulphate anion.).

Dose of the compound is 0.05 mg/kg - 1.0 mg/kg, based on 1 mg/kg of morphine. The cpd. is administered to the entrails. The cpd. is administered to outside of enteron with an injection, within two hours before the administration of the anaesthetic analgesic. The cpd. (I) is

methyl naltolexone.

USE/ADVANTAGE - For prevention and redn. of nausea and emesis caused from the use of morphine. (Provisional Basic previously advised in week 8916)

Abstract (Equivalent): US 4861781 A

Pharmaceutical compsn. contains one or more quat.
nor-hydroxymorphinone derivs. of formula (I), dispersed with the usual carriers and opt. additives. In (I), R is allyl, chloroallyl, propargyl or cyclopropylmethyl; and X is an anion, pref. halide or methosulphate.

USE- Cpds. (I) are used in dosages about 0.05-0.25~mg/kg to prevent or relieve nausea and emesis arising from the